

CLAIMS

1. (Currently Amended) A method of treatment, comprising:
identifying a human patient that is susceptible to ischemia; and
reducing the likelihood of an occurrence of a harmful effect of ischemia by
administering an effective amount of a stable free radical prior to the onset of ischemia;
wherein the likelihood is reduced in comparison to a human patient that was not
subjected to the administering step.
2. (Original) The method of Claim 1, wherein the nitroxide is 4-hydroxy-2,2,6,6-tetramethylpiperidine-1-oxyl.
3. (Original) The method of Claim 1, wherein the human patient's susceptibility to
ischemia arises from a medical procedure associated with a significant ischemic risk.
4. (Original) The method of Claim 3, wherein the medical procedure is the
treatment of a hemorrhage.
5. (Original) The method of Claim 3, wherein the medical procedure is the
treatment of an aneurysm.
6. (Currently Amended) The method of Claim 3, wherein the medical procedure is
surgery.
7. (Currently Amended) The method of Claim 3, wherein the medical procedure is
an endovascular procedure.
8. (Original) The method of Claim 1, wherein the mode of nitroxide administration
is selected from the group consisting of oral and intravenous administration.
9. (Currently amended) A method of treatment comprising:
identifying a patient scheduled to undergo a medical procedure involving a risk of
ischemia;
reducing the likelihood of an occurrence of a harmful effect of ischemia by
administering to the patient, prior to the medical procedure, an effective amount of a
stable free radical nitroxide;
performing the medical procedure; and
administering to the patient, an additional amount of a stable free radical nitroxide
to ameliorate a harmful effect of ischemia.

10. (Original) The method of Claim 9, wherein the nitroxide is 4-hydroxy-2,2,6,6-tetramethylpiperidine-1-oxyl.

11. (Original) The method of Claim 9, wherein the medical procedure is the treatment of a hemorrhage.

12. (Original) The method of Claim 9, wherein the medical procedure is the treatment of an aneurysm.

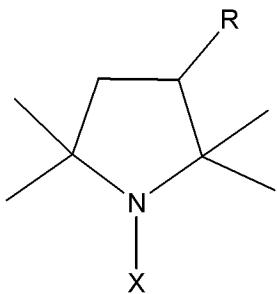
13. (Original) The method of Claim 9, wherein the medical procedure is surgery.

14. (Original) The method of Claim 9, wherein the medical procedure is an endovascular procedure.

15. (Original) The method of Claim 9, wherein the mode of nitroxide administration is selected from the group consisting of oral and intravenous administration.

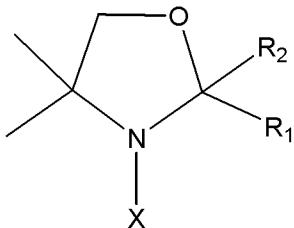
16-31 (Canceled)

32. (Currently Amended) The method of Claim 1 wherein the nitroxide is selected from the group consisting of



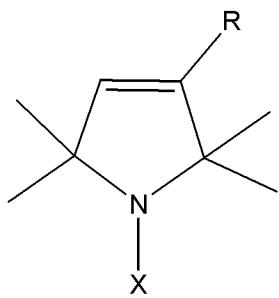
or a pharmaceutically acceptable salt thereof

wherein X is selected from O[•] and OH, and R is selected from COOH, CONH, CN, and CH₂ NH₂;



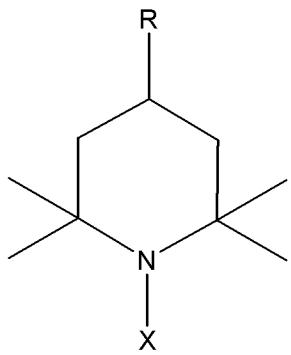
or a pharmaceutically acceptable salt thereof

wherein X is selected from O[•] and OH, and R₁ is selected from CH₃ and spirocylohexyl, and R₂ is selected from C₂ H₅ and spirocyclohexyl;



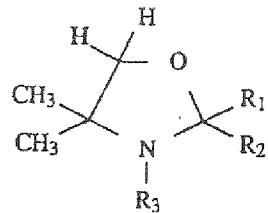
or a pharmaceutically acceptable salt thereof

wherein X is selected from O[•] and OH and R is CONH;

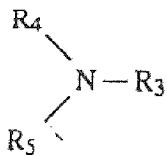


or a pharmaceutically acceptable salt thereof

wherein X is selected from O[•] and OH and R is H, OH, and NH₂;



wherein R₁ is -CH₃; R₂ is -C₂H₅; -C₃H₇; -C₄H₉; -C₅H₁₁; -C₆H₁₃; -CH₂-CH(CH₃)₂; -CHCH₃C₂H₅; or -(CH₂)₇-CH₃; or wherein R₁ and R₂ together form spirocyclopentane, spirocyclohexane, spirocycloheptane, spirocyclooctane, 5-cholestane, or norbornane; R₃ is -O[•] or -OH, or a physiologically acceptable salt thereof which has antioxidant activity;



wherein R_3 is $-O\cdot$ or $-OH$; and

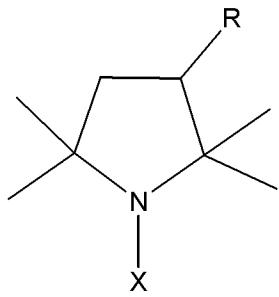
wherein R_4 and R_5 combine together with the nitrogen to form a heterocyclic group; wherein the atoms in the heterocyclic group (other than the N atom shown in the formula) may be all C atoms or may be C atoms and one or more N, O and/or S atoms; or

wherein R_4 and R_5 combine together to form substituted or unsubstituted pyrrole, imidazole, oxazole, thiazole, pyrazole, 3-pyrroline, pyrrolidine, pyridine, pyrimidine, or purine; or

wherein R_4 and R_5 themselves comprise a substituted or unsubstituted cyclic or heterocyclic group;

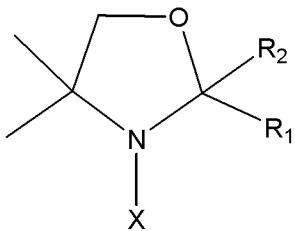
2-ethyl-2,5,5-trimethyl-3-oxazolidine-1-oxyl, 2,2,6,6-tetramethylpiperidine-1-oxyl (TEMPO), 4-hydroxy-2,2,6,6-tetramethylpiperidine-1-oxyl (TEMPOL), 4-amino-2,2,6,6-tetramethyl-1-piperidinyloxy (Tempamine), 3-Aminomethyl-PROXYL, 3-Cyano-PROXYL, 3-Carbamoyl-PROXYL, 3-Carboxy-PROXYL, 4-oxo-TEMPO, 4-amino-TEMPO, 4-(2-bromoacetamido)-TEMPO, 4-(ethoxyfluorophosphonyloxy)-TEMPO, 4-hydroxy-TEMPO, 4-(2-iodoacetamido)-TEMPO, 4-isothiocyanato-TEMPO, 4-maleimido-TEMPO, 4-(4-nitrobenzoyloxy)-TEMPO, and 4-phosphonoxy-TEMPO.

33. (Previously Presented) The method of Claim 9 wherein the nitroxide is selected from the group consisting of



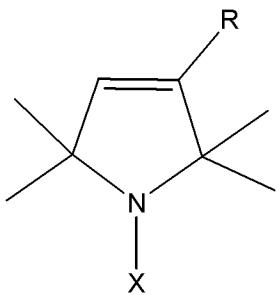
or a pharmaceutically acceptable salt thereof

wherein X is selected from O• and OH, and R is selected from COOH, CONH, CN, and CH₂ NH₂;



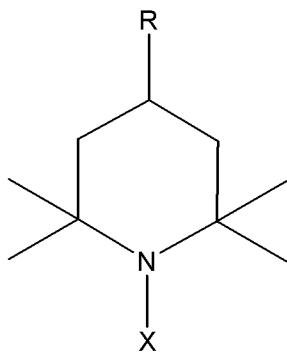
or a pharmaceutically acceptable salt thereof

wherein X is selected from O• and OH, and R₁ is selected from CH₃ and spirocyclohexyl, and R₂ is selected from C₂H₅ and spirocyclohexyl;



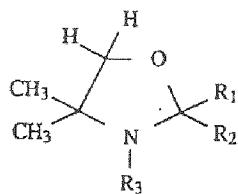
or a pharmaceutically acceptable salt thereof

wherein X is selected from O• and OH and R is CONH;

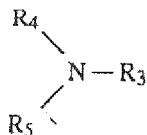


or a pharmaceutically acceptable salt thereof

wherein X is selected from O• and OH and R is selected from H, OH, and NH₂;



wherein R_1 is $-CH_3$; R_2 is $-C_2H_5$, $-C_3H_7$, $-C_4H_9$, $-C_5H_{11}$, $-C_6H_{13}$, $-CH_2-CH(CH_3)_2$, $-CHCH_3C_2H_5$, or $-(CH_2)_7-CH_3$, or wherein R_1 and R_2 together form spirocyclopentane, spirocyclohexane, spirocycloheptane, spirocyclooctane, 5-cholestane, or norbornane; R_3 is $-O\cdot$ or $-OH$, or a physiologically acceptable salt thereof which has antioxidant activity;



wherein R_3 is $-O\cdot$ or $-OH$; and

wherein R_4 and R_5 combine together with the nitrogen to form a heterocyclic group; wherein the atoms in the heterocyclic group (other than the N atom shown in the formula) may be all C atoms or may be C atoms and one or more N, O and/or S atoms; or

wherein R_4 and R_5 combine together to form substituted or unsubstituted pyrrole, imidazole, oxazole, thiazole, pyrazole, 3-pyrroline, pyrrolidine, pyridine, pyrimidine, or purine; or

wherein R_4 and R_5 themselves comprise a substituted or unsubstituted cyclic or heterocyclic group;

2-ethyl-2,5,5-trimethyl-3-oxazolidine-1-oxyl, 2,2,6,6-tetramethylpiperidine-1-oxyl (TEMPO), 4-hydroxy-2,2,6,6-tetramethylpiperidine-1-oxyl (TEMPOL), 4-amino-2,2,6,6-tetramethyl-1-piperidinyloxy (Tempamine), 3-Aminomethyl-PROXYL, 3-Cyano-PROXYL, 3-Carbamoyl-PROXYL, 3-Carboxy-PROXYL, 4-oxo-TEMPO, 4-amino-TEMPO, 4-(2-bromoacetamido)-TEMPO, 4-(ethoxyfluorophosphonyloxy)-TEMPO, 4-hydroxy-TEMPO, 4-(2-iodoacetamido)-TEMPO, 4-isothiocyanato-TEMPO, 4-maleimido-TEMPO, 4-(4-nitrobenzoyloxy)-TEMPO, and 4-phosphonoxy-TEMPO.

34. (Currently amended) A method of treatment comprising:

identifying a human patient who is susceptible to ischemia associated with a medical procedure; and

reducing a harmful effect of ischemia in the human patient after the medical procedure by administering an effective amount of 4-hydroxy-2,2,6,6-tetramethylpiperidine-1-oxyl prior to the onset of ischemia and prior to the medical procedure.

35. (Canceled) The method of Claim 34, wherein the nitroxide is 4-hydroxy-2,2,6,6-tetramethylpiperidine-1-oxyl.

36. (Previously Presented) The method of Claim 34, wherein the human patient's susceptibility to ischemia arises from a medical procedure associated with a significant ischemic risk.

37. (Currently Amended) The method of Claim 34, wherein the medical procedure is the treatment of a hemorrhage.

38. (Currently Amended) The method of Claim 34, wherein the medical procedure is the treatment of an aneurysm.

39. (Currently Amended) The method of Claim 34, wherein the medical procedure is surgery.

40. (Currently Amended) The method of Claim 34, wherein the medical procedure is an endovascular procedure.

41. (Previously Presented) The method of Claim 34, wherein the mode of nitroxide administration is selected from the group consisting of oral and intravenous administration.

42. (Currently Amended) A method of treatment comprising:

identifying a patient scheduled to undergo a medical procedure involving a significant risk of ischemia;

reducing a harmful effect of ischemia in the human patient after the medical procedure by administering an effective amount of 4-hydroxy-2,2,6,6-tetramethylpiperidine-1-oxyl performing the medical procedure; and

administering to the patient after the performing step, an additional amount of 4-hydroxy-2,2,6,6-tetramethylpiperidine-1-oxyl effective to reduce a harmful effect of ischemia

43. (Canceled) The method of Claim 42, wherein the nitroxide is 4-hydroxy-2,2,6,6-tetramethylpiperidine-1-oxyl.

44. (Previously Presented) The method of Claim 42, wherein the medical procedure is the treatment of a hemorrhage.

45. (Previously Presented) The method of Claim 42, wherein the medical procedure is the treatment of an aneurysm.

46. (Previously Presented) The method of Claim 42, wherein the medical procedure is surgery.

47. (Previously Presented) The method of Claim 42, wherein the medical procedure is an endovascular procedure.

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